

KOCHNEV, F.P., prof., doktor tekhn.nauk

Selecting directions for passenger train routes. Zhel.dor.transp.
41 no.3:55-56 № '59. (MIRA 12:6)
(Railroads--Passenger traffic)

KOCHNEV, F., prof., doktor tekhn. nauk

Foreword. Trudy NIIT no.116:3-4 '59.

(MIRA 12:11)

1. Nachal'nik Moskovskogo instituta inzhenerov shesnodorozhnogo
transporta im. I.V. Stalina,
(Railroads--Labor productivity)

KOCHNEV, F.P., prof., doktor tekhn.nauk

New trends in scientific, research, and educational institutes,
structural design bureaus and laboratories. Elek.i tepl.ting
4 no.2:28 F '60. (MIRA 13:6)

1. Nachal'nik Moskovskogo instituta inzhenerov sbelesnodorozhnogo
transporta.
(Electric engineering--Study and teaching)

KOCHNEV, F.P., prof., doktor tekhn.nauk

Determining the detention time of cars in classification yards.
Zhel.dor.transp. 42 no.6:84-89 Je '60. (MIRA 13:7)
(Railroads--Freight cars)

KOCHNEV, Fedor Petrovich, doktor tekhn.nauk, prof.; MAKSIMOVICH, Boris
Mikhailovich, kand.tekhn.nauk, dotsent; SOTNIKOV, Isaak
Bentsionovich, kand.tekhn.nauk, dotsent; SIMONOV, K.S.,
kand.tekhn.nauk, dotsent; MANYUKOV, O.S., inzh., red.;
BOEROVA, Ye.N., tekhn.red.

[Problems concerning the organisation of train movement] Voprosy
organisatsii dvizheniya poezdov. Moskva, Vses.iadatel'ske-poligr.
ob'edineniye M-va putei soobshcheniya, 1961. 211 p.

(MIRA 14:6)

(Railroads—Traffic)
(Railroads—Signaling)

KOCHNEV, Y.P., doktor tekhn.nauk, prof.

Some potentials of labor productivity in transportation. Trudy
MIIT no.137:169-178 '61. (MIRA 15:1)
(Railroads--Labor productivity)

KOCHNEV, F.P., doktor tekhn.nauk, prof.

Expediency of using rail motorears in suburban transportation. Elek.
i topl.tiaga 3 no.4:39-42 Ap '61. (MIRA 14:6)
(Railroad motorears)

KOCHNEV, F.P., prof., doktor tekhn.nauk

Effectiveness of diesel trains and rail motorcars in suburban
and local communications. Zhel.dor.transp. 43 no.2:37-40 F '61.
(Railroads—Commuting traffic) (Electric railroads)

KOCHNEV, Fedor Petrovich, prof., doktor tekhn.nauk; FUZIN, A.I.,
kand.tekhn.nauk, retsentsent; MANYUKOV, G.S., inzh., red.;
VOROTNIKOVA, L.F., tekhn. red.

[Organizing the transportation of suburban passengers in rail-
road motorcars] Organizatsiia peravevovok prigorodnykh passazhirov
avtomotrisami. Moskva, Transzheldorizdat, 1962. 153 p.
(MIRA 15:7)

(Railroad motorcars)

KOCHNEV, F.P., prof., doktor tekhn. nauk, zaslushennyi deyatel'
~~nauki i tekhniki~~

Locomotives and rolling stock used in commuter service should
be better utilized. Elek. i topl. tizn 7 no.3:22-24 M- '63.

(MIRA 16:6)

(Railroads—Electrification)
(Railroads—Passenger traffic)

KOCHNEV, P. P., prof., doktor tekhn. nauk, saslusheyny deyatel'
nauki i tekhniki RSFSR

Potentials of suburban train sheets. Zhel. dor. transp. 45
no.1:38-41 Ja '63. (MIRA 16:4)

(Railroads—Commuting traffic)

KOCHNEV, F.P., prof., doktor tekhn.nauk, zasluzhennyy deyatel' nauki i tekhniki RSPSR

Efficient methods for increasing the speed of passenger trains in local communications. Zhel.dor.transp. 45 no.9:42-45 3 '63.

(MIRA 16:9)

(Railroads--Train speed)

KOCHNEV, F.P., doktor tekhn. nauk, prof.

Problems of the reduction of the costs of construction of new
railroads. Trudy MIIT no.162:3 '63. (MIRA 17:4)

1. Rektor Moskovskogo instituta inzhenerov zheleznodorozhnogo
transporta.

KOCHNEV, P.P., prof., doktor tekhn. nauk, red.

[Use of electronic digital computers in calculations connected with the operation of railroads.] Primenenie elektronnykh tsifrovyykh vychislitel'nykh mashin v raschetakh, svyazannykh s ekspluatatsiei zheleznykh dorog. Moskva, Transport, 1965. 221 p. (Trudy Moskovskogo instituta inzhenerov zheleznodorozhnogo transporta, no.202). (MIRA 19:6)

1. Rektor Moskovskogo instituta inzhenerov zheleznodorozhnogo transporta.

KOCHNEV, Fedor Petrovich, zasl. deyatel' nauki i tekhniki RSFSR
doktor tekhn. nauk prof.; DLUGACH, B.A., red.

[Weight and speed of passenger trains] Voz i skorost'
passazhirekikh poezdov. Moskva, Transport, 1965. 275 p.
(MIRA 19:1)

KOCHNEV, I. (Bodaybo, Irkutskaya obl.); ISHMAYEV, I. (Bodaybo, Irkutskaya obl.)

Under the conditions of northern regions. Posh. delo 8
no.9:25 8 '62. (MIRA 16:11)

SILEROVA, A.I.; KACHNEV, I.K.

Changes in infrared absorption spectra of the brain tissue
of a white rat dehydrated by vacuum drying. Vest. LGU 19
no.16:38-41 '64.

(MIRA 17:11)

PODINEV, L.; VASII'YEV, M.; FILATOV, S.

Neutralization of exhaust gases. Avt.transp. 40 no.1:19-21 Ja
'62. (Auto 15:1)
(Automobile exhaust gas)

GALE, M. V.

"Air Conditioning in Mining of Ore Deposits. Dis. sent to Scientific Commission."
Sub 23 Mar 51, Inst of Mining, Acad Sci USSR.

Dissertations presented for science and engineering degrees in Moscow during 1951.

CC: Sub. No. 400, 2 May 55.

KOCHNEV, K. V.

KOCHNEV, K. V. -- "Control of Firedamp in Mining Deposits Susceptible to Self-Ignition." Sub 27 Jun 52, Inst of Mining, Acad Sci USSR (Dissertation for the Degree of Doctor in Technical Sciences)

SO: Vechernaya Moskva, January-December 1952

KOCHNEV, K. V.

USSR/Mining - Underground Fires, Mine Safety

Jul 52

"Prevention of Endogenous Fires in Copper -Pyrite Mines," A. A. Ivanov, K. M. Charkviani, N. P. Diyev, K. V. Kochnev, Z. G. Sheina, Ye. F. Jordan, F. N. Pavlov

"Iz Ak Nauk, Otdel Tekh Nauk" No 7, pp 1037-1044

Presents results of works conducted since 1947 by a group of Soviet investigators studying causes of underground fires and establishing preventive measures. Discusses selection of mining system safe in respect to fires, sifting as basic preventive measures against underground fires, and ventilation for cooling ore rocks and for maintaining normal temp conditions in mines. Submitted by Acad A. A. Skochinskiy 1 Apr 52.

PA 228T99

KOCHNEV, K.V.

Problem of fire damp leakage through caved ground in copper
pyrite mines. Trudy Gor.-geol.inst. no.27:122-133 '55.

(MLRA 9:9)

(Mine ventilation)

KOCHNEV, K.V.

Computing mine ventilation in heated untouched ore blocks.

Trudy Gor.-geol.inst. no.27:134-139 '55.

(MLRA 9:9)

(Mine ventilation)

KOCHNEV, K.V., doktor tekhnicheskikh nauk.

Main trends in technical engineering studies on silicosis prevention in enterprises in the Urals. Sbor. rab. po sil. no.1:3-11 '56. (MLRA 10:2)

1. Gorno-Geologicheskii institut Ural'skogo filiala Akademii nauk SSSR.

(LUNGS--DUST DISEASES) (URALS--MINE SANITATION)

KOCANEV, K.V., doktor tekhnicheskikh nauk.

Technical means of preventing silicosis in Urals' mines. Gor.shur.
no.2:63-64 F '56. (MLRA 9:5)

1. Gorno-geologicheskii institut Ural'skogo filiala AN SSSR.
(Ural Mountain region--Minners' phthisis) (Ural Mountain region
--Mine sanitation)

BUTAKOV, Sergey Yefimovich; TALIYEV, V.N., kand.tekhn.nauk, retsenzent;
KOCHNEV, K.Y., doktor tekhn.nauk, red.; DUGINA, N.A., tekhn.red.

[Air ducts and ventilators; aerodynamics of ventilator equipment]
Vozdukhoprovody i ventilatory; aerodinamika ventilatornykh
ustanovok. Moskva, Gos.nauchno-tekhn.isd-vo mashinostroit. lit-ry,
1958. 35 p. (MIRA 11:6)
(Ventilation)

KOCHNEV, K.V.

Industrial prevention of silicosis in the U.S.S.R. and in
foreign countries. Trudy Gor.-geol. inst. UFAI SSSR no.31:
125-172 '58. (MIRA 12:9)
(MINE DUSTS) (MINERS--DISEASES AND HYGIENE)

KOCHNEV, K.Y.; FILATOV, S.S.

Aerodynamics of flow in open-cut mining. Trudy Gorn.-geol. inst.
UZAN SSSR no.31:245-250 '58. (MIRA 12:9)
(Strip mining) (Mine ventilation)

KOCHNEV, K.V.

Air conditioning in modern mines. Trudy Gor.-geol.inst.UFAN
SSSR no.41:35-46 '59. (MIRA 13:5)
(Mine ventilation) (Air conditioning)

KOCHNEV, Konstantin Vasil'yevich; BUTAKOV, S.Ye., otv.red.; MAKOVSKIY,
O.M., red.isd-va; ASTAF'YEVA, O.I., tekhn.red.

[Temperature control in mines] Teplovoe konditsionirovanie
rudnichnogo vozdukh. Moskva, Izd-vo Akad.nauk SSSR, 1960.
133 p. (Akademiya nauk SSSR. Ural'skii filial, Sverdlovsk.
Gorno-geologicheskii institut. Trudy, no.44) (MIRA 13:9)
(Mine ventilation)

KOCHNEV, K.Y., prof., doktor tekhn.nauk; FILATOV, S.S., mladshiy nauchnyy
sotrudnik

Improvement of atmospheric conditions in deep quarries. Sbor. rab.
po silik. no.2:3-14 '60. (MIRA' 14:3)

1. Gorno-geologicheskiy institut Ural'skogo filiala AN SSSR.
(MINE VENTILATION) (MINE DUSTS)

KOCHNEV, K.V., prof.doktor tekhn.nauk; YEFREMOVA, T.K., gornyy inzhener

Controlling dust pollution of the air in mechanised longwall mining
in the Chelyabinsk Coal Basin. Sbor. rab. po silik. no.2:125-129
'60. (RIRA 14:3)

1. Gorno-geologicheskii institut Ural'skogo filiala AN SSSR.
(CHELYABINSK BASIN—DUST—PREVENTION)

FILATOV, S.S.; KOCHNEV, K.V.; VASIL'YEV, M.V.

Searching for practical methods of controlling exhaust gases from
truck haulage in strip mines. Gor.zhur. no.5:65-66 Ny '60.
(MIRA 14:3)

1. Ural'skiy filial AN SSSR, Sverdlovsk.
(Mine sanitation) (Automobile exhaust gas)

D'YAKOV, Vasil'y Vasil'yevich; KOCHNEV, K.V., otv. red.; GRISHAYENKO,
M.I., red. izd-va; SHKLYAR, S.Ye., tekhn. red.

[Dust control in scraper levels] Obespylivanie gorizontov akre-
peroveniia. Moskva, Gos. nauchno-tekhn. izd-vo lit-ry po gor-
nomu delu, 1961. 90 p. (MIRA 14:5)
(Mine dusts)

KOCHNEV, K.V., prof., doktor tekhn.nauk

Results of scientific studies on the technical prevention of silicosis.
Sbor. rab. po silik. no3:3-9 '61, (MIRA 15:10)

1. Gorno-geologicheskiy institut Ural'skogo filiala AN SSSR.
(Mine dusts) (Lungs—Dust diseases)

KOCHNEV, K.V., prof., doktor tekhn.nauk; REZNIKOV, N.A., gornyy inzh.; FILATOV,
S.S., gornyy inzh.

Controlling dust formation in the Korkino open-pit mine.
Sbor. rab. po silik. no.3:79-85 '61.

(MIRA 15:10)

1. Gorno-geologicheskiy institut Ural'skogo filiala AN SSSR, trest
Korkinugil'.

(Chelyabinsk Basin—Coal mines and mining) (Mine dusts)

KOCHNEV, K.V., prof., doktor tekhn.nauk; SHEINA, Z.G., kand.khimicheskikh nauk;
FEDOROVA, G.G., kand.khimicheskikh nauk

Preventing dust formation and keeping down floating dust in the
Korkino open-pit mine. Sbor. rab. po silik. no.3:109-117 '61.
(MIRA 15:10)

1. Gorno-geologicheskii institut Ural'skogo filiala AN SSSR.
(Chelyabinsk basin--Mine dusts)

KOCHNEV, K.V., prof., doktor tekhn.nauk; D'YAKOV, V.V., gornyy inzh.; KOVALEV,
V.I., gornyy inzh.

Effect of the speed and initial dust content of an air current on its
picking up dust from the surface of mine workings. Sbor. rab. po
sil'm. no.3:119-128 '61. (MIRA 15:10)

1. Gorno-geologicheskii institut Ural'skogo filiala AN SSSR i Sverdlovskiy
institut okhrany truda.

(Mine dusts)

KOCHNEV, K.V., prof., doktor tekhn.nauk; SHEINA, Z.G., kand.khim.nauk;
FEDOROVA, G.G., kand.khim.nauk

Wetting agents and saline additives as means of controlling the
process of dust prevention. Bor'ba s sil. 5:21-27 '62.
(MIRA 16:5)

1. Gorno-geologicheskiy institut Ural'skogo filiala AN SSSR.
(Mine dusts—Prevention)

KORNIY, K.V., prof. doktor tekhn. nauk; U.S.S.R., U.S.S.R., kand. tekhn. nauk;
U.S.S.R., U.S.S.R., kand. tekhn. nauk

Testing of electric mine filters. B.I.R. s. 11. 6164-167 '64
(MIRA 1812)

KOCHNEV, M.

33206. KOCHNEV, M. Trud i presnya, -dva pokoleniya (ocherki ob ivan. tekstil'schikakh ivanovskiy al'ranakh, kn. 11, 1949, c. 1953

SO: Letopis' Zhurnal'nykh Statey, Vol. 45, Moskva 1949

KOCHNEV, M

W/5
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.27K7

Zinaida Men'shikova. Moskva, Moskovskiy rabochiy, 1951

62 p. port.

KOCHNEV, Mikhail

The fifth year of the seven-year plan. Av. i korm. 45 no.1:2-5
Ja '63. (MIRA 16:1)

(Russia—Economic policy)

KOCHNEV, M.F.; GALYGIN, A.V.

Designing continuous pickling units. Metallurg 10
no.1:31-32 Ja '65.

(MIRA 18:4)

1. Nachal'nik listoprokatnogo tsekh Magnitogorskogo
metallurgicheskogo kombinata (for Kochnev). 2. Nachal'nik
travil'nogo otdeleniya Magnitogorskogo metallurgicheskogo
kombinata (for Galygin).

137-58-6-12155

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 6, p 140 (USSR)

AUTHORS: Benyakovskiy, M.A., Shadrin, V.A., Kulikov, V.I.,
Uzivenko, A.M., Kustobayev, G.G., Kochnev, M.F.,
Kutuyev, Ya.S.

TITLE: The Interrelation of the Pressure, the Pull, and the Thickness
of a Strip Subjected to Cold Rolling (Vzaimosvyaz' davleniya,
natyazheniya i tolshchiny lenty pri kholodnoy prokatke)

PERIODICAL: Byul. nauchno-tekhn. inform. Ural'skiy n.-i. in-t chernykh
metallov, 1957, Nr 3, pp 114-123

ABSTRACT: A three-stand rolling mill of the MMK was employed during
research concerned with the effect of rolling (R) rate on the
thickness of a strip (S), the establishment of interrelation of
pressure and pull during cold R, and determination of the sig-
nificance of longitudinal and transverse thickness variations in
the S. A mathematical relationship is established between the
basic parameters of the technological process of cold R of a S.
It is established that variations in the tension of the strip mid-
way between the stands of a mill have a decisive effect on the
formation and magnitude of thickness variations in the S.

Card 1/2

137-58-6-12155

The Interrelation of the Pressure, the Pull, and the Thickness of a Strip (cont.)

Fluctuations of R rate at the MMK have practically no effect on the thickness of the S. Variations in the pull produce thickness variations in the S equivalent to 0.01-0.02 mm on the average.

S.N.

1. Steel--Processing 2. Steel--Pressure distribution 3. Rolling mills--Applications

Card 2/2

500-NEV, M. P.
BENTAKOVSKIY, M.A.; KULIKOV, V.I.; SHADRIN, V.A.; KOLPAKOV, I.P.; KUTUYEV,
Ya.S.; KUSTOMAYEV, G.G.; KOCHNEV, M.P.; YESIFOV, I.V.; PETROV, B.I.

Power consumption for the deformation of metal and conditions of
strip rollings. Stal' 17 no.1:59-63 Ja '57. (NIRA 10:3)

1. Ural'skiy institut chernykh metallov i Magnitogorskiy metallur-
gicheskiy kombinat.

(Rolling (Metalwork))

GUSEV, Vasily Fedorovich; KVITKO, M.M., inzh., retsentsent; KOCHNEV,
M.G., inzh., retsentsent; BASNETSYAN, A.A., inzh., red.;
GORDENKOVA, L.P., tekhn.red.

[Assembly and adjustment of vertical propeller pumps] Montazh
i naladka vertikal'nykh propellernykh nasonov. Moskva, Gos.
nauchno-tekhn.isd-vo mashinostroit.lit-ry, 1960. 197 p.
(MIRA 13:11)

(Pumping machinery)

TATARINOV, M.P., prof.; KOCHNEV, M.G., inzh.; CHESNOKOV, A.V., inzh.

New centrifugal mine pump. Nauch. trudy MOI no.23:209-216
'58. (MIRA 15:12)

(Mine pumps)

RABINOVICH, N.I.; KUNDZICH, M.M., insh., retsentsant; KOCHMEV, M.O.,
insh., red.; ZHURAVLEVA, M.N., red.i:d-va; TIKHANOV, A.Ya.,
tekhn. red.

[Mobile pumping stations for irrigation] Peredvishnye nasosnye
stantsii dlia orosheniia. Moskva, Mashgis, 1962. 115 p.

(MIRA 16:3)

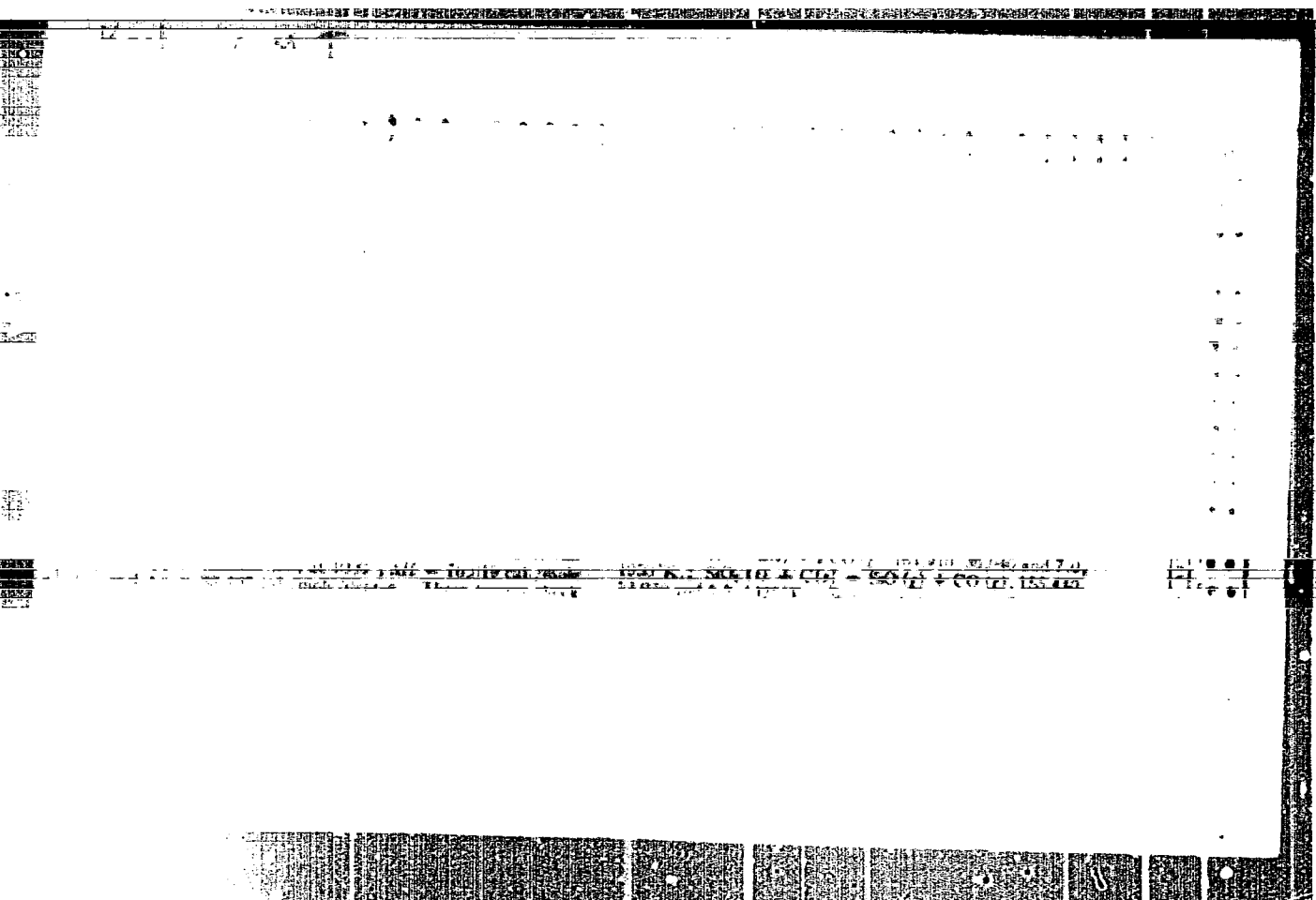
(Pumping stations) (Irrigation)

CA

Characterization of dust from reactors of Krasnoarsk Works. P. I. Ulyashin and A. I. Kabanov. *Tr. Vsesoyuzn. Nauch. Issled. Inst. Khim. Akad. Nauk SSSR*, 1968, No. 7, 71-75. (Phys. and Chem. Anal.)
 Analyses were made of the dust at various points of the dust-collector system of the Krasnoarsk Copper Works. The approx. compn. of the charge was: concentrate 10%, quartz 20, return 20, limestone 12, quartz with sulfide (30% CuS) 20 tons. The mass carried by the gas from the Cottrell precipitators is a material of - 225 mesh. The sp. gr. of the dust varies from 3.61 to 3.71, and the analysis is: Cu 7.0-8.97, Zn 6.0-7.25, Fe 21.2-23.3 and 10.0-10.2%. The Cu and Zn content in the dust is higher than their content in the waste mass; the Fe content, however, is highest in the coarsest fractions. The Fe content is about the same in all sizes. The following compn. of Cu were found: Cu_2O , Cu_2S , Cu_2S , Cu_2S . The Cu_2S is the predominant Cu compound and constitutes 40% of the Cottrell dust. The sulfate and oxide of Cu increase, and the sulfide content decreases in the direction of the gas motion. Commercial oxidation takes place on the Cottrell precipitators; the sulfides are oxidized better than oxides and sulfates. B. M. Standish

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KOCHNOV, M. I.

M. I. Kochnov, Examination of abnormal sectors of the logarithmic curves of the dissociation pressure of cobalt arsenides. P. 1210

This work is devoted to the study of the dissociation pressure of cobalt arsenides. This investigation was preceded by the study of ignition temperatures of cobalt arsenides by heating them in a stream of air. This work is to augment the characteristics of arsenides by studying their behavior in a neutral atmosphere near their ignition temperature and on the given dissociation pressure to characterize the cobalt arsenides thermodynamically.

Chair of Metallurgy of Heavy Non-Ferrous Metals of the Ural Industrial Institute, March 22, 1948

SO: Journal of Applied Chemistry (USSR) 21, No. 12 (1948)

KOCHNEV, M. I.

PA 11/49T15

USSR/Chemistry - Silicon Oxide
Chemistry - Vapor Pressure

Aug 48

"Vapor Tension of Silicon Oxide," P. V. Gel'd,
M. I. Kochnev, Ural Ind Inst imeni S. M. Kirov, 4 pp

"Dok Ak Nauk SSSR" Vol LXXI, No 4

Studies vapor pressure of silicon dioxide between
900° and 1100° C. Tabulates results. Independence
of SiO vapor tension of its degree of volatilization
shows that solid state is a separated phase, and not
a solid solution. Submitted 15 May 48.

11/49T15

CA

Free-energy and heat-content changes in the reaction of formation of cobalt arsenide. *J. L. Kice, J. Chem. Phys.* 19, 716-7 (1951). (1) Ignition temps. and ΔH vapor pressures (at a temp. close to the ignition temp.), in mm. Hg, were found for the following arsenides: CoAs , 840° , 2.3×10^{-3} (840°); CoAs , 810° , 1.45×10^{-3} (810°); CoAs , 770° , 9.5×10^{-4} (770°); CoAs , 730° , 6.97×10^{-4} (730°); CoAs , 690° , 5.81×10^{-4} (730°). (2) For the formation reaction $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, from the vapor pressure, the free energy of formation at 730° is $\Delta F = -17,477 \text{ cal./mole}$, the heat between 730° and 700° , $\Delta H = -41,540 \text{ cal./mole}$, and the entropy change at 730° , $\Delta S = -22.18 \text{ cal./mole/deg}$. The change of the dissociation equl. const. $K_p = p_{\text{As}_2}^{-1/2}$ (atm.), between 730° and 700° , is satisfactorily rendered by $\log K_p = (3942/T) - 4.840$. For fused CoAs , the dissociation vapor pressure between 980° and 1080° is rendered by $\log p (\text{atm.}) = (13694/T) - 9.502$; hence, for the formation at 980° , $\Delta F = -17,867$, and $\Delta H = -39,810 \text{ cal./mole}$. Actually, the products of dissociation are partially dissolved in the original CoAs , resulting in a liquid phase of variable composition; consequently, the vapor-pressure formula has only an approx. validity. (3) For the reaction $3 \text{CoAs (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 737° , $\Delta F = -1981 \text{ cal.} = -1724 \text{ cal./mole CoAs}$, and $\Delta H = -20,064 \text{ cal.} = -4123 \text{ cal./mole CoAs}$; at the melting temp. of CoAs , 940° , $\Delta F = -6280 \text{ cal.} = -1298 \text{ cal./mole CoAs}$, and $\Delta H = -15,105 \text{ cal.} = -3023 \text{ cal./mole CoAs}$. Between 730° and 737° , $\log K_p = (4334/T) - 2.482$. From the data for the foregoing 3 formation reactions, it is calculated for $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 737° , $\Delta F = -1913$, $\Delta H = -39,895 \text{ cal./mole}$. (4) For $3 \text{CoAs (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 770° , $\Delta F = -8483 \text{ cal.} = -4241 \text{ cal./mole CoAs}$, $\Delta H = -21,208 \text{ cal.} = -10,604 \text{ cal./mole CoAs}$; at the melting temp. of CoAs , 944° , $\Delta F = -6171 \text{ cal.} = -3085 \text{ cal./mole CoAs}$, $\Delta H = -11,940 \text{ cal./mole CoAs}$. Between 730° and 770° , $\log K_p = (3922/T) - 2.677$; between 730° and 944° , $\log K_p = (2947/T) - 1.948$, and between 944° and 980° , $\log K_p = (2680/T) - 1.268$. Hence, for $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 770° , $\Delta F = -17,820$, $\Delta H = -41,600 \text{ cal./mole Co}_3\text{As}_2$. (5) For $\text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 730° , $\Delta F = -5564 \text{ cal.} = -2782 \text{ cal./mole CoAs}$, $\Delta H = -30,707 \text{ cal.} = -15,353 \text{ cal./mole CoAs}$; in the temp. ranges 740° - 760° , 790° - 820° , and 890° - 1000° , $\log K_p = (4420/T) - 2.804$, $(6782/T) - 2.768$, and $(6944/T) - 2.558$, resp. For the formation from the elements, $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 730° , $\Delta F = -18974$, $\Delta H = -39,790 \text{ cal./mole Co}_3\text{As}_2$. (6) For $3 \text{CoAs (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 616° , $\Delta F = -1967$, $\Delta H = -15,855 \text{ cal./mole CoAs}$; between 601° and 740° , $\log K_p = (3466/T) - 2.474$. For the formation from the elements, $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 740° , $\Delta F = -22,285$, $\Delta H = -39,548 \text{ cal./mole Co}_3\text{As}_2$. (7) For $\text{CoAs (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 840° , $\Delta F = -4945 \text{ cal.} = -2473 \text{ cal./mole CoAs}$.

7) = 2.482. From the data for the foregoing 3 formation reactions, it is calculated for $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 737° , $\Delta F = -1913$, $\Delta H = -39,895 \text{ cal./mole}$. (4) For $3 \text{CoAs (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 770° , $\Delta F = -8483 \text{ cal.} = -4241 \text{ cal./mole CoAs}$, $\Delta H = -21,208 \text{ cal.} = -10,604 \text{ cal./mole CoAs}$; at the melting temp. of CoAs , 944° , $\Delta F = -6171 \text{ cal.} = -3085 \text{ cal./mole CoAs}$, $\Delta H = -11,940 \text{ cal./mole CoAs}$. Between 730° and 770° , $\log K_p = (3922/T) - 2.677$; between 730° and 944° , $\log K_p = (2947/T) - 1.948$, and between 944° and 980° , $\log K_p = (2680/T) - 1.268$. Hence, for $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 770° , $\Delta F = -17,820$, $\Delta H = -41,600 \text{ cal./mole Co}_3\text{As}_2$. (5) For $\text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 730° , $\Delta F = -5564 \text{ cal.} = -2782 \text{ cal./mole CoAs}$, $\Delta H = -30,707 \text{ cal.} = -15,353 \text{ cal./mole CoAs}$; in the temp. ranges 740° - 760° , 790° - 820° , and 890° - 1000° , $\log K_p = (4420/T) - 2.804$, $(6782/T) - 2.768$, and $(6944/T) - 2.558$, resp. For the formation from the elements, $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 730° , $\Delta F = -18974$, $\Delta H = -39,790 \text{ cal./mole Co}_3\text{As}_2$. (6) For $3 \text{CoAs (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 616° , $\Delta F = -1967$, $\Delta H = -15,855 \text{ cal./mole CoAs}$; between 601° and 740° , $\log K_p = (3466/T) - 2.474$. For the formation from the elements, $3 \text{Co (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow \text{Co}_3\text{As}_2 (\text{cryst.})$, at 740° , $\Delta F = -22,285$, $\Delta H = -39,548 \text{ cal./mole Co}_3\text{As}_2$. (7) For $\text{CoAs (cryst.)} + \frac{1}{2} \text{As}_2 (\text{gas}) \rightarrow 3 \text{CoAs (cryst.)}$, $K_p = p_{\text{As}_2}^{-1/2}$, at 840° , $\Delta F = -4945 \text{ cal.} = -2473 \text{ cal./mole CoAs}$.

ΔH_f of Co_2As = -14,790 cal. = -8075 cal./mole Co_2As ;
 between 600 and 620° and 640-660°, $\log K_p$ = (2128/7)
 -1.267 and (2128/7) - 2.268, resp. Discuss. pressure
 - calcd. by the last equation in the last temp. range coincides
 with data obtained by eq. solution, and agree with the
 pressure of 700.0 mm. Hg. ded. by heating Co_2As in a
 stream of H_2 at 675°. For the formation from the ele-
 ments, $\text{Co (cryst.)} + \frac{1}{2} \text{As (gas)} \rightarrow \text{Co}_2\text{As (cryst.)}$, at
 700°, and ΔH_f = -12,500, ΔH_f = -27,260 cal./mole
 Co_2As . (7) Values of ΔF and ΔH of formation from the
 elements, at 700°, per 1 g.-atom Co, are: Co_2As , -2720
 and -2920; Co_3As , -6000 and -10,314; Co_4As , -8115
 and -12,600; Co_5As , -9200 and -20,700; Co_6As ,
 -11,300 and -20,120; Co_7As , -12,500 and -27,260
 cal./Co; in the same order, ΔF and ΔH per 1 g.-atom As,
 are: -6000 and -10,314; -8115 and -20,700; -9175
 and -20,120; -9200 and -20,700; -7440 and
 -10,410; -6000 and -10,700 cal./As. At 1014°, the
 values are, per 1 g.-atom Co, for Co_2As , -2020 and
 -7200; Co_3As , -5000 and -9120; Co_4As , -6000 and
 -12,620 cal./Co. In the same order, per 1 g.-atom As,
 -7200 and -10,200; -7070 and -10,204; -7200 and
 -10,210 cal./As. Thus, at the same temp., ΔF and ΔH
 per 1 g.-atom Co increase regularly from lower to higher
 N. Then

KOCHNEV, M. I.

Methods of preparing arsenides of easy composition. M. I. Kochnev (J. Appl. Chem. USSR, 1959, 23, 561). Abstract: Arsenides of As are obtained from pure As₂O₃ by reduction with charcoal in an atm. of H₂. Co is obtained by reduction of Co₂O₃ with H₂ and Ni or Fe by reduction of its oxide. Electrolytic Cu is used. The powdered metal and As are mixed in stoichiometric proportions in a porcelain crucible. The crucible is placed inside a horizontal quartz tube inside a tubular electric furnace. Air is displaced by H₂ and the temp. is kept above the m.p. of the arsenide. Inside the quartz tube As vapours concentrate in the lower part, while the upper part of the tube is filled with H₂. The molten metal reacts quickly with vapours of As and the whole synthesis is completed within 1 hr. Co₂As, Co₃As, Co₂As₂, CoAs, Ni₂As, Ni₃As, Ni₂As₂, Co₂As₂, and Fe₂As obtained in this way do not differ from their theoretical composition by more than 0.05%. To obtain higher arsenides it is necessary to maintain the temp. above the m.p. of As in hermetically closed, steel, pressure vessels, lined with a mixture obtained from Fe₂O₃, Al₂O₃, and Fe powder wetted with water-glass and calcined at 850°. The products obtained in pressure cylinders have an As content slightly higher than theoretical; this excess can be removed by heating in vac. at 400-500° in an ampoule with a small opening. J. H. J. Zana.

BC

A-1
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Methods of preparing emulsion of definite composition
M. I. Kucharsky. Applied Chem. U.S.S.R. 22, 202 (1951)
(1951) (Eng. Translation).—See C.A. 44, 7007d R. M. S.

1952

KOCHNEV, M.I.

①

The Thermodynamic Properties of the Cobalt Azides.
M. I. Kochnev (Doklady Akad. Nauk S.S.S.R., 1960, 78, (3), 433-435). (In Russian). The thermal dissociation of the Co azides was studied by measuring the pressure of the liberated As vapour by the method of flow. Synthetic CoAs_2 , Co_2As_2 , Co_3As_2 , Co_4As_2 , and Co_5As_2 were studied, and equations for their dissociation are given. Temp. of experiment ranging from 400° to 1000° C. were used, depending on the temp. of the azide. The mol. of gaseous azide was assumed to be As_2 , and as the dissociation pressure of the azides in the given temp. interval was very low, the ideal gas laws were assumed to be applicable. The dissociation pressures of the six Co azides at 57° C. from their ignition temp. are given, and the equilibrium const. of the dissociation reactions are calculated from them. The heats of formation of the Co azides and their heats of dissociation are also calculated. Solubility of the dissociation products in the Co azides affects the results slightly. Empirical equations for the dependence of the equilibrium const. on temp. are given.—Z. S. B.

[Handwritten signature: S. B. 2/14/64]

KUCHREV, M. I.

✓ Solid solutions of the system cobalt-arsenic. M. I. Kuchrev. Doklady Akad. Nauk S.S.S.R. 93, 1187 (1957).
 Alloys of Co-As in the range of 44-70% Co were investigated by the method of ignition temps. Plots of ignition temps. vs. % As yield 4 peaks corresponding to the arsenides, each with a higher ignition temp. as the % As increases. The expected horizontal temp. lines between the arsenides and the stepwise rise does not appear. The temp. drops abruptly from each max., becomes horizontal and then rises to the ignition temp. of the subsequent arsenide. This is explained on the assumption that solid solns. of limited soly. are formed; the following approx. compn. is given:

$\text{CoAs} + 48.4\% \text{ Co}$; $\text{Co}_2\text{As}_3 + 51-58.3\% \text{ Co}$; $\text{Co}_3\text{As}_4 + 58.5-61.8\% \text{ Co}$; $\text{Co}_4\text{As}_5 + 61.8-67.5\% \text{ Co}$. I. B.

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Dissociation Pressure of Copper Arsenide. M. I. Koshov (Doklady Akad. Nauk S.S.S.R., 1961, 78, (3), 563-565). (In Russian). The dissociation pressure (P) of an alloy containing 71-80% Cu, 20-30% As (Cu_3As contains 28.2% As) was measured over the range 500°-700° C. by the molecular-flow method used previously for metal arsenides (Zhur. Prikl. Khim., 1945, 18, 1210; preceding column). Over the range 500°-600° C., $\log P \propto 1/T$, where T is the abs. temp., but in the interval 600°-695° C. P drops sharply, and at 600° C. it could not be measured. Beginning at 605° C., however, P again increases with temp., so that at 695° C. it has reached the value it had at 500° C. By 700° C., P has begun to fall gradually. Comparison of the $\log P$ vs. $1/T$ curve with the equilibrium diagram shows that anomalies in the curve correspond to temp. of isothermal transformations in alloys with compn. near to Cu_3As . Over the temp. range 500°-600° C., the equilibrium const. K_p for the formation of Cu_3As from the elements was given by: $\log K_p = 11.90 - 1.25$. The free energy of formation at 500° C. $\Delta F_{500} = -6472$ cal./g.-mol. For the interval 500°-600° C., the change in heat content $\Delta H = -11,303$ cal./g.-mol. and the change in entropy $\Delta S = -8.7$ cal./g.-mol. The value of 26,000 cal./g.-mol. for the heat of formation obtained by Raveleberg (Metall u. Erz, 1936, 14, 279) is criticized. G. V. K. T.

KOCHNEV, K. I.

262T23

USSR/Metallurgy - Nickel Ores, Smelting Aug 52

"Effect of the Oxides of Alkali Metals on the Viscosity of High-Magnesium Slags," N. P. Diyeu, M. I. Kochnev, M. F. Kochin, Yu. A. Gyrdymov

Iz Ak Nauk SSSR, OTN, No 8, pp 1182-1188

Discusses exptl smelting of Ni ores for obtaining complete characteristic of compn and properties of slags. Establishes that slags with 10-12% FeO and 17-19% MgO are extremely difficult to melt and have high viscosity. Presence of alkali metal oxides in amount of 1.5-2.0% reduces high fusibility and viscosity of these slags, promoting their removal

262T23

from furnace. Such favorable effect of oxides takes place only when total content of silica and alumina does not surpass approx 52%. Presented by Acad I. P. Bardin 28 Oct 51.

The structure of copper scale and the mechanism for the oxidation of copper. V. E. Dey and M. I. Kichnev. Doklady Akad. Nauk S.S.S.R. 85, 461-463 (1972). Based on the literature data, a mechanism is proposed for the oxidation of Cu based on the diffusion of O through the crystal lattice. The thickness of the Cu₂O layer increases in two directions: on the boundary Cu-Cu₂O and on the boundary Cu₂O-Cu₂CO₃. J. Rostov Leach.

KOCHNEV, M.I.; ZAYIDMAN, T.I.

Characteristics of variation in the physico-chemical properties
of copper selenide. Izv. AN SSSR Otd. tekhn. nauk no. 12:1813-1818
D '53. (MLRA 7:2)

1. Predstavleno akademikom I.P. Bardinym. (Copper compounds)

KOCHETOV, N. I.

Chemical Abstracts
May 25, 1954
General and Physical
Chemistry

Oxidation of selenium-copper alloys. N. P. Kochetov, M. I. Kochetov, A. P. Ivanova, and I. N. Zakharenko. *Tr. Akad. Khim. Nauk SSSR* (1953) 2187. The kinetics of oxidation of Cu alloys containing 0, 0.44, 1.77, and 2.47% Se was investigated. The rate of oxidation between 600 and 1000° increases with Se content, although between 800° and 900° the rate is lower than for pure Cu. The rate-time curves are parabolic. After 1.5 hrs. at 1000°, Se content drops from 2.47 to 2.27% in the metal and to 0.050% in the oxide scale. Adding Ag, as in 1.37% Se-1.24% Ag triple alloy, increases the rate of oxidation. This is of interest in the refining of Cu. E. M. Ellis.

"APPROVED FOR RELEASE: 09/18/2001

CIA-RDP86-00513R000723520010-1

APPROVED FOR RELEASE: 09/18/2001

CIA-RDP86-00513R000723520010-1"

KOCHNEV, M.I.

DIYEV, Nikolay Pavlovich; VERMENICHYV, Sergey Aleksandrovich; PEN'KO, Aleksandr Stepanovich. KOCHNEV, M.I., redaktor; LUCHKO, Yu.V., redaktor; KOVALENKO, N.I., tekhnicheskii redaktor

[Smelting copper ores and concentrates in reverberatory furnaces]
Otrashatel'naya plavka mednykh rud i koncentratov; uchebnoe posobie
dlia shkol i kursov masterov. Sverdlovsk, Gos. nauchno-tekhn. izd-vo
lit-ry po chernoi i tsvetnoi metallurgii, 1954. 300 p. (MLBA 8:4)
(Copper--Metallurgy) (Smelting furnaces)

KOCHNEV, M. I.

AID - P-86

Subject : USSR/Chemistry

Card : 1/1

Authors : Diyev, N. P., Kochnev, M. I., Paduchev, V. V., and Sloridse, G. Ya.

Title : Formation of manganese sulfate from manganese dioxide by ferrous sulfate or pyrite

Periodical : Zhur. Prikl. Khim. 27, no. 4, 356-359, 1954

Abstract : Roasting of manganese dioxide with ferrous sulfate at 700-800°C converts 92-99% manganese into sulfate; with pyrite, 87.5-89% Mn. Three references (U.S.S.R.): 1935-1944. Three graphs, three tables.

Institution : None

Submitted : February 16, 1953

~~Characterization of iron oxide dissociation. M. I. Kochnev, E. V. Oshin, D. A. Kozlov, and N. S. Smirnov. Zhur. Prikl. Khim. 1954, No. 4, 163-7. Ref. 1. Zhur. Khim. 1954, No. 4, 356-359. — Partial pressure of O (P_O) of Fe₂O₃ was detd. at 800-1350°. In the region of small pressures (up to 1100°) the method of root efflux was used, in which samples of chemically pure Fe₂O₃ were placed in quartz ampuls and were degassed for 3-4 hrs. in vacuo at 600-800°. For temps. above 1150° P_O was detd. monometrically, porcelain app. being used. In the interval 1100-1420° log P_O = (-32,405/T) + 6.10 and for temps. of 920-1100° log P_O = (-14,852/T) + 4.80. The thermal effects of the reaction for the above are 145,270 and 68,684 cal./mole. At temps. 800 and 850-900° deviations from the above relations were observed; at these temps. P_O exceeds the calcd. values 100-300 times. The points of anomalous behavior correspond to transformation of Fe₂O₃. At 931° the red modification of Fe₂O₃ changes to black and at 817° a Fe₂O₃ changes to γ-Fe₂O₃.~~

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Kochner, M I.

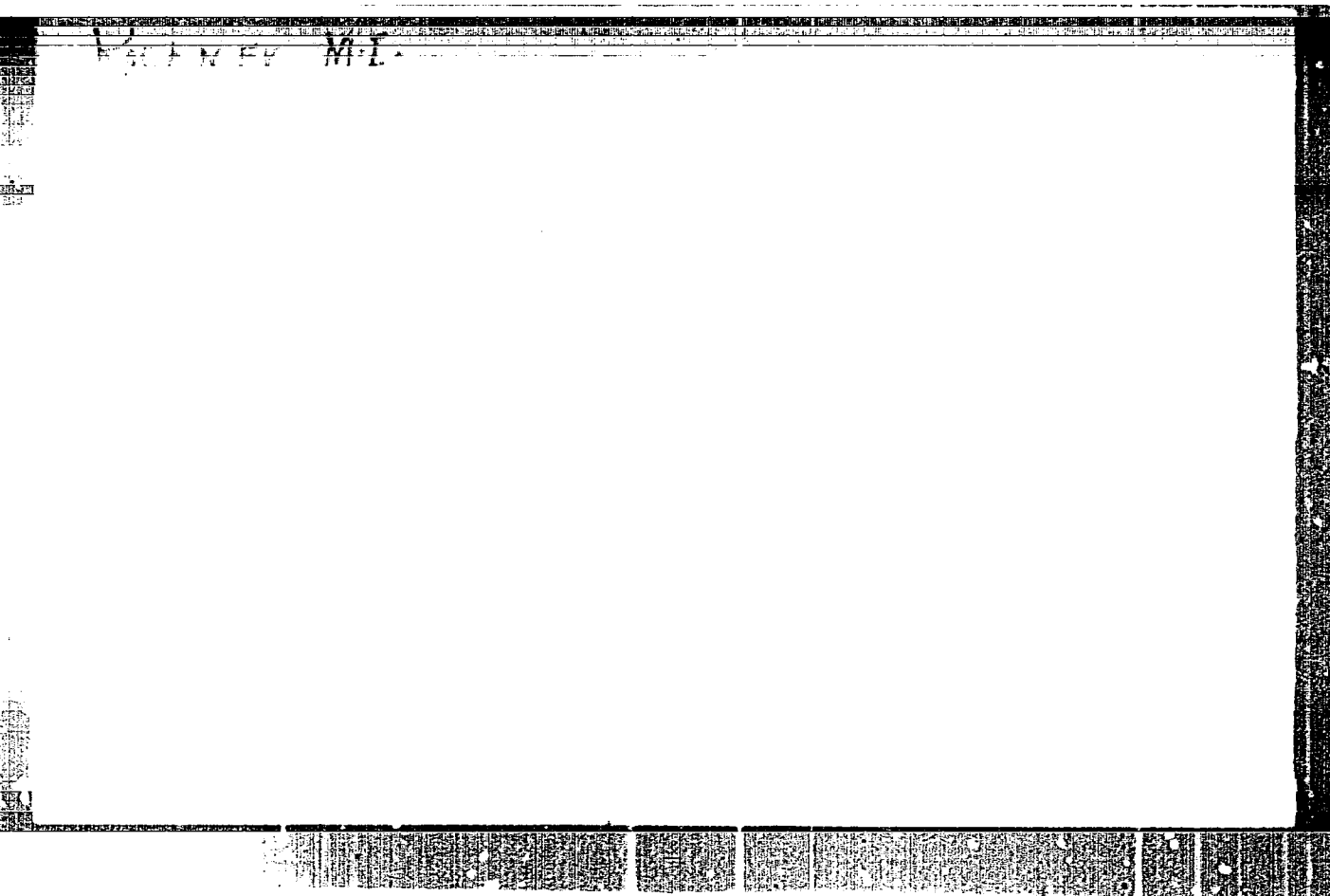
✓ The relation between the changes in dissociation pressure and the electrical resistance of arsenic. M. I. Kochner. Doklady Akad. Nauk SSSR, 195, 1958, 116-117. The quant. relation between the elec. and physical therm. properties of semiconductors, such as Ge , was studied by a comparison of vapor pressure and the elec. cond. between 700 and 800°. The curves are similar in shape, with 2 max. near 64° and 700°. The 64° max. is related to the transformation in the Ge structure at 64°. The 700° max. is explained by the transformation of the cubic lattice into the hexagonal 2 conditions in W. M. I.

AGLITSKIY, Viktor Aleksandrovich; KOCHENY, M.I., redaktor; **KEL'NIK, V.P.,**
redaktor izdatel'stva; **KOVALENKO, N.I.,** tekhnicheskiy redaktor

[The manufacture of copper wire bars; theory and practice]
Proizvodstvo mednykh valerbarsov; teoriia i praktika. Sverdlovsk,
Gos. nauchno-tekhn. izd-vo lit-ry po cherno i tsvetnoi metallurgii,
Sverdlovskoe otd-nie, 1956. 287 p. (MLR 9:10)
(Wire) (Copper--Metallurgy)

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CIA-RDP86-00513R000723520010-1"

USSR/ Cosmochemistry. Geochemistry. Hydrochemistry

D.

Abs Jour : Referat Zhur - Khimiya, No 4, 1957, 11520

Author : Zaydman T.W., Kochnev M.I., Flekhanov A.P.

Inst : Department of Technical Sciences, Academy of Sciences USSR

Title : On Reactivity and Thermal Transformations of Sphalerite

Orig Pub : Izv. AN SSSR, Otd. tekhn. n., 1956, No 6, 168-171

Abstract : Investigated was the dependence of heating rate of a specimen of sphalerite on the temperature at a constant flow of heat. Chemical composition of specimen (in %): Zn 65.15, S 32.25, Pb 0.42, Fe 0.78. Correlation between rate of heating (degree/minute and temperature is represented by a graph which is of the nature of sudden changes with a gradual subsidence on increase in temperature to 650°. The curve shows two sharply manifested maxima (at 650 and 950°) and two minima (at 850 and 1020°), after which the curve extends upward. Anomalous progression course of the graph is due to the occurrence of a number of thermal transformations associated with changes in state of crystal lattice. A correlation has been ascertained between intervals of thermal transformations of ZnS and temperature of critical points of the forming elements.

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VOLKOVA, P.I.; DITNY, N.P.; KOCHNEV, M.I.

Phase composition of copper-zinc mattes. Izvet. met. 29 no.1:
47-50 Ja '56. (MIRA 9:6)
(Copper--Metallurgy) (Zinc--Metallurgy)

SOV/137-58-10-21303

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 10, p 123 (USSR)

AUTHORS: Zalazinskiy, G. G., Diyev, N. P., Kochnev, M. I.

TITLE: On the Kinetics and the Mechanism of Oxidation of a Copper-nickel Alloy at Elevated Temperatures (O kinetike i mekhanizme okisleniya medno-nikelevogo splava pri vysokikh temperaturakh)

PERIODICAL: Tr. in-ta metallurgii. Ural'skiy fil. AN SSSR, 1957, Nr 1, pp 59-64

ABSTRACT: The kinetics of oxidation (O) of a Cu-Ni alloy (I) in the process of multiple heating and the diffusion of O_2 in I were investigated, and comparative data on the oxidation in air and in O_2 were determined. Attempts to establish selective diffusion of Cu and Ni in the process of 30 hours' O of I containing 4.1% Ni were unsuccessful. The rate of diffusion of O_2 in I in the process of its O at 1000°C in air is higher than the rate of the growth of the scale, as the result of which I becomes saturated with O_2 . The rate of O of I quickly increases with the saturation of its entire mass with O_2 . The rate of O of I in either air or O_2 in the $700 - 1000^\circ$ temperature range increases - 2.2 times with a 100° increase in temperature. The rate of O of I in O_2 is on

Card 1/2

SOV/137-58-10-21303

On the Kinetics and the Mechanism of Oxidation (cont.)

the average 1.56 times higher than in air. The rate of O of I in air at 700 - 800° is lower than the rate of O of electrolytic Cu, whereas at 900 - 1000° it is higher than that of pure Cu.

N. L.

1. Copper-nickel alloys--Oxidation
2. Copper-nickel alloys--Temperature factors
3. Copper--Diffusion
4. Nickel--Diffusion

Card 2/2

Translation from: Referativnyy zhurnal. Metallurgiya, 1958, N- 12, p 53 (USSR) SOV/137-58-12-24303

AUTHORS: Volkova, P. I., Diyev, N. P., Kochnev, M. I.

TITLE: The Behavior of Zinc Compounds When Matte is Allowed to Stand
(Povedeniye soyedineniy tsinka pri otstaivani shleyznov)

PERIODICAL: Tr. In-ta metallurgii. Ural'skiy fil. AN SSSR. 1957, Nr 1, pp 87-92

ABSTRACT: A Kryptol furnace and alundum crucibles are used under laboratory conditions to run experiments in which Cu mattes containing ~25% Cu and various amounts of Zn are allowed to stand. It is established that up to ~8% of the ZnS in a melt of Cu matte at 1140°C is in solution. If the ZnS content is higher, the excess is in the form of a ZnS-enriched sphalerite solid solution, which rises to the surface when the matte bath is allowed to stand and forms a thick viscous mass that complicates separation of the matte and the slag. The $ZnOFe_2O_3 + FeS = ZnS + Fe_3O_4$ reaction occurring in the melts causes the bulk of the Zn ferrite to convert to ZnS when allowed to stand for a long period. The magnetite is concentrated by segregation in the lower portion of the matte ingot, and also in its upper portion, which

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The Behavior of Zinc Compounds When Matte is Allowed to Stand (cont.) SOV/137-58-12-24303

is of elevated viscosity when allowed to stand. Small amounts of SiO_2 and Al_2O_3 found in matte melts do not precipitate.

Ye. Z.

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SOV/137-58-12-24300

Translation from: Referativnyy zhurnal. Metallurgiya, 1958, Nr 12, p 53 (USSR)

AUTHORS: Volkova, P. I., Diyeu, N. P., Kochnev, M. I.

TITLE: Reaction Between Zinc Matte and Metallic Iron (Vzaimodeystviye mezhdu tsinkovistym shteynom i metallicheskim zhelezom)

PERIODICAL: Tr. In-ta metallurgii. Ural'skiy fil. AN SSSR, 1957, Nr 1, pp 93-98

ABSTRACT: The reaction between ZnS and metallic Fe in the 600-1300°C temperature range is studied. The thermodynamic calculation of the isobar potential is studied. It shows that the probability that the reaction would occur rises with temperature. A study is made of the rate of reaction between Zn matte and metallic Fe. It is established that: 1) The quantity of Zn removed in the process at a given Fe content rises with temperature; 2) the reaction between the solid phases occurs in the first 20 min, after which it ceases; 3) when the Fe content in the specimen is low and the temperature is 1000°, the reaction between Zn matte and Fe hardly occurs at all; 4) the marked change in the isobar potential of the reaction between ZnS and metallic Fe at 900° corresponds to the maximum Zn removal

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Reaction Between Zinc Matte and Metallic Iron
established by kinetic investigations.

L. S.

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SOV/137-58-7-14554

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 7, p 87 (USSR)

AUTHORS: Skorniyakov, G.P., Motova, Z.A., Kochnev, M.I.

TITLE: Luminosity of Converter Flame During the Fining of Blister Copper (O svecenii plameni konvertera v period dovodki chernovoy medi)

PERIODICAL: Byul. tsvetn. metallurgii. 1957, Nr 24, pp 20-22

ABSTRACT: A photometric study was conducted of the converter flame (by spectroscope connected to a film camera), and optical monitoring by photoelectric pyrometer with recording potentiometer was performed. In both cases, intensity of emission was found to increase during the fining period, this being connected with an increase in the Cu contents of the melt. Maximum emission of light was attained during the final 5 to 8 min. The subsequent sharp drop is apparently to be explained by the onset of Cu oxidation.

1. Flames--Spectographic analysis 2. Photometry--Appli- L.P.
cations 3. Photoelectric pyrometers--Applications 4. Copper
--Properties

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Kochnev, M.I.

24-58-3-9/38

AUTHORS: Kochnev, M.I., Plotnikova, A.F. and Starkov, L.N. (Sverdlovsk).

TITLE: Temperature Features of the Process of Oxidation of Copper Sulphide (Temperaturnyye osobennosti protsessa okisleniya sul'fida medi)

PERIODICAL: Izvestiya Akademii Nauk SSSR, Otdeleniye Tekhnicheskikh Nauk, 1958, Nr 3, pp 82-88 (USSR)

ABSTRACT: Modern conceptions on the changes in the electron structure of atoms and their influence on the character of the chemical bond forces, gained from the study of semi-conductors, justify a more thorough analysis of various phenomena involved in metallurgical processes. In earlier work in this field, relating to compounds of heavy non-ferrous metals, the team of the authors of this paper established the existence of a temperature correspondence in the changes of the properties of these compounds and of their components. The aim of the work described in this paper was to study the interrelation between the temperature changes and the properties of copper and sulphur and the properties of the simple compound Cu_2S formed from these. Cu_2S was chosen for investigation due to the fact that it is one of the basic components of the raw materials for which new processes of roasting and smelting are being developed. Two specimens were investigat-

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24-58-3-9/38

Temperature Features of the Process of Oxidation of Copper Sulphide.

ed which were prepared synthetically by smelting, the respective compositions being the following: 78.9% Cu, 21.1% S and 79.1% Cu, 20.9% S (the theoretical composition being 79.8% Cu, 20.2% S). In both cases the composition was in the range of solid solutions of sulphur in Cu_2S . The differences in the composition of the two specimens proved to be of little importance. The experiments were carried out mainly with sulphide grains of the sizes 0.50 to 0.63 mm. Oxidation of the sulphide was carried out in a vacuum set-up, a sketch of which is shown in Fig.1, p.83, using the method of circulating air in a closed system, drawing it through a layer of the charge which is heated to a certain temperature; the gases obtained after drawing off the air through the charge were caught by a device in which cooling by means of liquid nitrogen was applied for the purpose of freezing out sulphurous acid anhydride and sulphuric anhydride. According to Averbukh, B.D. (Ref.7) the quantity of forming sulphuric anhydride under these conditions is very low and, therefore, was not determined separately. The investigations were carried out with a constant initial air pressure in the system

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Temperature Features of the Process of Oxidation of Copper Sulphide.

of 408 mm and a constant temperature of the charge, which were established during each measurement of oxygen consumption after three minutes. The total duration of the experiment was 21 or 30 mins. The graph, Fig.2, gives the change in the speed of oxidation of Cu_2S during continuous heating. The graph, Fig.3, gives the temperature dependence of the coefficient of electrical resistance of the copper. The graph, Fig.4, gives the temperature dependence of the oxidation speed of copper sulphide along a fresh surface. The graph, Fig.5, gives the temperature dependence of the quantity of copper which is present in the form of oxides and sulphate in the residue on the degree of oxidation of Cu_2S .

In Fig.6 the consumption of oxygen and the yield of sulphurous acid anhydride as a function of the temperature are graphed. It was established that the speed of oxidation of Cu_2S does not change continuously with temperature but is

complicated by a number of anomalous deviations within narrow temperature ranges. The temperatures of the narrow deviations in the kinetics of oxidation of copper sulphide are critical temperatures for pure copper and pure sulphur, the manifestation of which is considered as being the result

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Temperature Features of the Process of Oxidation of Copper Sulphide.

of changes in the electron structures of the atoms with increasing temperature. The character of the changes in the speeds of oxidation of the copper sulphide at critical temperatures is elucidated and the limits of anomalous temperature ranges were determined; in most cases the sharp fluctuations in the oxidation speed reach 20 to 30% and take place in the temperature range 5 to 15°C. On the basis of study of the oxidation isotherms, the degree of utilisation of the oxygen and the yield of sulphurous acid anhydride and of the influence of the oxidation duration, the conclusion is arrived at that the determining factor in the process of oxidation of Cu_2S at temperatures up to 450°C is the formation of sulphate. There are 6 figures and 12 references, 11 of which are Soviet, 1 English.

ASSOCIATION: Institut metallurgii Ural'skogo filiala, AN SSSR
(Institute of Metallurgy, Ural Branch Ac.Sc., USSR)

SUBMITTED: January 3, 1957.

Card 4/4 1. Copper sulfide--Oxidation 2. Temperature--Effects

[illegible]

SOV/24-58-4-22/39

AUTHORS: Kochnev, M.I. and Plotnikova, A.F. (Sverdlovsk)

TITLE: Kinetics of Reducing Magnetite at Temperatures at Which Transformation of Iron Takes Place (Kinetika vosstanovleniya magnitnogo zheleznyaka pri temperaturakh prevrashcheniy zheleza)

PERIODICAL: Izvestiya Akademii Nauk SSSR, Otdeleniye Tekhnicheskikh Nauk, 1958, Nr 4, pp 118 - 121 (USSR)

ABSTRACT: In earlier work one of the authors of the paper found that there is a direct relation and a temperature correspondence between the changes taking place in various metal compounds and the changes in the properties of the individual elements forming these compounds. In this paper, the authors aimed at verifying this conclusion for iron oxides and thus to try and explain anomalous phenomena taking place in reduction processes. To bring the theoretical investigations nearer to industrially used materials, the authors used in the experiments magnetite and not pure oxides. The composition of the ore was as follows: 51.1% Fe, 22.9% FeO, 0.18% MnO, 12.0% MgO, 0.55% CaO, 1.74% Al_2O_3 , 9.4% SiO_2 , 2.6% S, 0.05% Cu. The kinetics of

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Kinetics of Reducing Magnetite at Temperatures at Which Transformation of Iron Takes Place

reduction were investigated on a circular vacuum test rig (described in an earlier paper) (Ref 20) inside a hydrogen atmosphere; the initial pressure was 408 mm, hydrogen was sucked through a layer 25 mm high, weighing 7 g, with particle dimensions between 0.4 and 0.6 mm. The temperature in the layer was measured with an accuracy of $\pm 1^\circ\text{C}$; the circulation speed was 600 ml./min with a volume of the system of 800 ml. The gaseous reaction products were frozen out in a trap which was cooled by liquid nitrogen. The ore was heated to the desired temperature in vacuum (10^{-2} to 10^{-3} mm Hg). Following that, a quantity of hydrogen was introduced which was equal in every case, maintaining the pressure constant. The speed of reduction was judged from the consumption of hydrogen in the closed system which was measured every minute. Every three minutes the system was joined to a vacuum and, after that, the temperature was readjusted and the gaseous phase renewed. It could, therefore, be assumed that a number of successive measurements, carried out at 3 min intervals, provided the isotherm of the reduction of the ore. At

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each temperature the experiments were carried out with a fresh ore specimen. Since the aim of the work was to elucidate the existence of a relation between the changes in the properties of the iron, at temperatures at which it is known that phase transformations take place, and the kinetics of reduction of iron oxides under equal conditions, the investigations were carried out in the temperature range 700 to 910 °C. In this temperature range magnetic as well as polymorphous transformations take place in the iron at 768 and 906-910 °C, respectively. The speeds of reduction of magnetite, expressed in terms of hydrogen consumption during the first 3 min as a function of the temperature, are graphed in Figure 1. It can be seen that the graph contains several anomalous sections. The obtained results justify revision of certain views expressed

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on the kinetics of reduction of iron oxides. In the first instance, it is quite evident that sintering phenomena, changes in the porosity and recrystallisation of the studied substances and in the reaction products, changes of speed of diffusion and the speed of chemical reactions do occur in the course of variation of the temperature during reduction of the ores and of iron oxides. The most plausible explanation of the anomalous phenomena in the processes of reduction is based on the changes of the state of the iron atoms as a function of the temperature, particularly as regards phase transformations. The obtained results (Figure 1) indicate that the jumps in the speed of reduction at 752-756 °C correspond to the magnetic transformation of the iron and not to the Curie point of the magnetite. Thus, the change in the character of the chemical process at the

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temperature of magnetic transformation of the iron confirms the conclusion that the anomalous phenomena in the chemical and physical processes at various temperatures are based on the changes in the state of the atoms and not on changes of the crystal lattice. It can be seen from the graph, Figure 2 (temperature coefficient of the electric resistance of iron) that the thermal coefficient of the electric resistance changes appreciably in the temperature range 400 to 450 °C and also at 550 and 650 °C.

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The following conclusions are arrived at:

- 1) The speed of reduction of magnetite with hydrogen decreases in jumps and then again increases within narrow temperature range approaching the following temperatures: magnetic transformation of the iron (752°C), $\text{Fe}_{\beta} \rightarrow \text{Fe}_{\gamma}$ transformation (906°C) and at about 820°C , at which the properties of the iron change considerably.
- 2) Using the example of the jump-like change in the speed of reduction at temperatures of the magnetic transformation of iron it is shown that the observed anomalies in the kinetics of the process are based on the changes of the electron structure of the iron atoms.
- 3) The anomalies in the kinetics of reduction of ore are explained more satisfactorily than hitherto from the point of view of transformations taking place in the iron.
- 4) The results of the here described work can be applied for selecting optimum temperatures for the preparation and processing of ores in the neighbourhood of critical points.

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Kinetics of Reducing Magnetite at Temperatures at Which
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There are 2 figures and 27 references, 2 of which are
Swedish, 1 German, 1 English and 23 Soviet.

ASSOCIATION: Institut metallurgii Ural'skogo filiala AN SSSR
(Institute of Metallurgy, Ural Branch of the
Ac.Sc.USSR)

SUBMITTED: March 11, 1957

Card 7/7

AUTHOR: Kochnev, M. I. (Sverdlovsk)

SOV/24-58-11-32/42

TITLE: On the Nature of Isothermal Processes in the Blast Furnace (O prirode izotermicheskikh protsessov v domennoy pechi)

PERIODICAL: Izvestiya Akademii Nauk SSSR, Otdeleniye Tekhnicheskikh Nauk, 1958, Nr 11, pp 121-123 (USSR)

ABSTRACT: In investigating the heat exchange processes in blast furnaces, the charge of which contains ores and fluxes, it was established that the curve of temperature distribution along the height of the furnace is S-shaped (Refs 1-8). It is characteristic that in the central part the temperature in the vertical direction changes very little and is maintained spontaneously in the neighbourhood of 800 to 900°C; the difference between the temperature of the gases and of the charge assumes in this region the minimum value of 5 to 10°C. The presence of this quasi isothermal zone, referred to as the no-load reserve zone or the zone of moderate temperatures, was considered as being a stable, typical relation. However, recent experiments by N. N. Babarykin and F. A. Yushin (Ref 9) have shown that if fluxed agglomerate is smelted without charging ore and limestone, the previously

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On the Nature of Isothermal Processes in the Blast Furnace

mentioned temperature distribution is not maintained and the temperature curves show two to three isothermal breaks. The cause of these has not been clarified. On the basis of data obtained from practical experience and analysis of the thermal balance of blast furnaces, it can be concluded that in the reserve zone of the blast furnace a considerable development of reduction processes takes place and also an intensification of the processes of heat exchange which is brought about by the isothermal processes. From the correspondence between the temperatures of the change in the properties of iron and its oxides and the temperatures at which the break in the temperature rise occurs in blast furnaces it can be concluded that the appearance of isothermal zones is due to transient states of iron and its oxides, i.e. it is due to the nature of the main reacting substances.

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SOV/24-58-11-32/42

On the Nature of Isothermal Processes in the Blast Furnace

There are 23 references, all of which are Soviet.

ASSOCIATION: Institut metallurgii Ural'skogo filiala AN SSSR
(Institute of Metallurgy, Ural Branch of the Ac.Sc. USSR)

SUBMITTED: April 17, 1957

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PLEKHANOV, A.F.; KOCHNEV, M.I.

Phase transformations of solid state zinc. Trudy Inst.met.UFAN
SSSR no.3:37-42 '59. (MIRA 13:4)
(Zinc) (Phase rule and equilibrium)